The listing of claims will replace all prior versions, and listings, of claims in the application: <u>Listing of Claims</u>:

1. (Currently Amended) A compound of formula (I) or a pharmaceutical or veterinary acceptable salt, hydrate or solvate thereof

$$Q \xrightarrow{R_1} Q \xrightarrow{R_2} Q \xrightarrow{R_3} R_3 \xrightarrow{P_4} Y \xrightarrow{A} \qquad (I)$$

wherein

Q represents a radical of formula -N(OH)CH(=O) or formula -C(=O)NH(OH);

Y represents -C(=O)-, -C(=S)-, -S(=O)-, or $-SO_2$ -;

 R_1 represents hydrogen, C_1 - C_6 alkyl or C_1 - C_6 alkyl substituted by one or more halogen atoms, or, except when Q is a radical of formula -N(OH)CH(=O), a hydroxy, C_1 - C_6 alkoxy, C_1 - C_6 alkenyloxy, halogen, amino, C_1 - C_6 alkylamino, or di- $(C_1$ - C_6 alkylamino group;

 R_2 represents a substituted or unsubstituted C_1 - C_6 alkyl, C_1 - C_3 alkyl-O- C_1 - C_3 alkyl, C_1 - C_3 alkyl-S- C_1 - C_3 alkyl, cycloalkyl(C_1 - C_3 alkyl)-, aryl(C_1 - C_3 alkyl)-, heterocyclyl(C_1 - C_3 alkyl)-, or R^1R^2N - C_1 - C_3 alkyl group wherein R^1 represents hydrogen or C_1 - C_3 alkyl and R^2 represents C_1 - C_3 alkyl, or R^1 R^2N -represents a cyclic amino group;

R₃ and R₅ independently represent hydrogen or a substituted or unsubstituted C₁-C₆ alkyl group or R₃ and R₅ taken together with the carbon and nitrogen atoms to which they are respectively attached form a saturated heterocyclic ring of from 5 to 7 ring atoms, which may be fused to a second carbocyclic or heterocyclic ring, either of which rings may optionally be substituted;

R₄ represent hydrogen or a substituted or unsubstituted C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆

alkynyl, cycloalkyl, aryl, heterocyclyl, C_1 - C_3 alkyl-O- C_1 - C_3 alkyl, C_1 - C_3 -alkyl-S- C_1 - C_3 alkyl)-, cycloalkyl(C_1 - C_3 alkyl)-, heterocyclic(C_1 - C_3 alkyl)- or aryl(C_1 - C_3 alkyl)- group; and

A represents a primary, secondary or tertiary amino group or a group- R_6 , -OR₆, wherein R_6 is a substituted or unsubstituted C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, cycloalkyl, aryl, heterocyclyl, C_1 - C_3 alkyl-O-(C_1 - C_3 alkyl-O-(C_1 - C_3 alkyl-S-(C_1 - C_3 alkyl-NH-(C_1 - C_3 alkyl)-, cycloalkyl(C_1 - C_3 alkyl)-, heterocyclic(C_1 - C_3 alkyl)-1 or aryl(C_1 - C_3 alkyl)- group.

- 2. (Original) A compound as claimed in claim 1 wherein Q is an N-formyl hydroxylamine group -N(OH)CH(=O).
- 3. (Currently Amended) A compound as claimed in claim 1 or claim 2-wherein -Y- is -C(=O)- or -SO₂-.
- 4. (Currently Amended) A compound as claimed in any of the preceding claims claim 1 wherein R₁ is hydrogen.
- 5. (Currently Amended) A compound as claimed in any of the preceding claims claim 1 wherein R_2 is

optionally substituted C₁-C₆ alkyl, C₃-C₆ alkenyl, C₃-C₆ alkynyl or cycloalkyl;

phenyl(C_1 - C_6 alkyl)-, phenyl(C_3 - C_6 alkenyl)- or phenyl(C_3 - C_6 alkynyl)- optionally substituted in the phenyl ring;

cycloalkyl(C_1 - C_6 alkyl)-, cycloalkyl(C_3 - C_6 alkenyl)- or cycloalkyl(C_3 - C_6 alkynyl)-optionally substituted in the cycloalkyl ring; or

 $CH_3(CH_2)_pO(CH_2)_q$ - or $CH_3(CH_2)_pS(CH_2)_q$ -, wherein p is 0, 1, 2 or 3 and q is 1, 2 or 3.

- 6. (Currently Amended) A compound as claimed in any of claims 1 to 4-claim 1 wherein R₂ is methyl, ethyl, n- or iso-propyl, n-or iso-butyl, n-pentyl, iso-pentyl, 3-methyl-but-1-yl, n-hexyl, n-heptyl, n-acetyl, n-octyl, methylsulfanylethyl, ethylsulfanylmethyl, 2-methoxyethyl, 2-ethoxyethyl, 3-hydroxypropyl, allyl, 3-phenylprop-3-en-1-yl, prop-2-yn-1-yl, 3-phenylprop-2-yn-1-yl, 3-(2-chlorophenyl)prop-2-yn-1-yl, but-2-yn-1-yl, cyclopentyl, cyclopentyl, cyclopentylmethyl, cyclopentylpropyl, acyclohexylmethyl, cyclopentylpropyl, acyclohexylmethyl, cyclohexylethyl, cyclohexylpropyl, furan-2-ylmethyl, furan-3-methyl, tetrahydrofuran-2-ylmethyl, piperidinylmethyl, pyrid-2-ylmethyl, pyrid-3-ylmethyl, pyrid-4-ylmethyl, phenylpropyl, 4-chlorophenylpropyl, 4-methylphenylpropyl, 4-methylphenylpropyl, 4-methoxybenzyl.
- 7. (Currently Amended) A compound as claimed in any of claims 1-to 4-claim 1 wherein R_2 is (C_1-C_6) alkyl-, cycloalkylmethyl-, (C_1-C_3) alkyl-S- (C_1-C_3) alkyl-, or (C_1-C_3) alkyl- (C_1-C_3) alkyl-.
- 8. (Original) A compound as claimed in claim 7 wherein R₂ is a n-propyl, n-butyl, n-pentyl, cyclopentylmethyl, cyclopexylmethyl or cyclohexylethyl.
- 9. (Currently Amended) A compound as claimed in any-of-the preceding claims claim 1 wherein R_4 is hydrogen, (C_1-C_6) alkyl-, cycloalkylmethyl-, (C_1-C_3) alkyl-S- (C_1-C_3) alkyl-, or (C_1-C_3) alkyl-O- (C_1-C_3) alkyl-.
- 10. (Original) A compound as claimed in claim 9 wherein R₄ is hydrogen, methyl, ethyl, n-propyl, n-butyl, n-pentyl, cyclopentylmethyl, cyclopentylethyl, cyclohexylmethyl or cyclohexylethyl.
- 11. (Currently Amended) A compound as claimed in any of the preceding claims claim 1 wherein R_3 and R_5 , when not part of a ring, are independently hydrogen, (C_1-C_6) alkyl-, cycloalkylmethyl-, (C_1-C_3) alkyl-S- (C_1-C_3) alkyl-, or (C_1-C_3) alkyl-O- (C_1-C_3) alkyl-.
- 12. (Original) A compound as claimed in claim 11 wherein R₃ and R₅ are independently hydrogen, methyl, ethyl, n-propyl, n-butyl, n-pentyl, cyclopentylmethyl, cyclopentylethyl, cyclohexylmethyl or cyclohexylethyl.

13. (Currently Amended) A compound as claimed in any of claims 1 to 10 claim 1 wherein R_3 and R_5 taken together with the carbon and nitrogen atoms to which they are respectively attached form the following rings, wherein any sulfur atom present as a ring member may be oxidized to -SO- or $-SO_2$ -, R_4 is as defined in any preceding claim 1, and R represents hydrogen or C_1 - C_4 alkyl:

14. (Currently Amended) A compound as claimed in any of the preceding claims claim 1 wherein A is a group -NR₆R₇, -R₆, or -OR₆ wherein R₆ and R₇ independently represent a radical of formula (IV)

$$\frac{1}{2} (Alk^1)_m - (X)_p - (Alk^2)_n - Z \qquad (IV)$$

wherein

m, p and n are independently 0 or 1;

Z represents hydrogen or a carbocyclic or heterocyclic ring of 5 to 7 ring atoms which is optionally fused to a saturated or unsaturated carbocyclic or heterocyclic second ring of 5 to 7

ring atoms;

Alk¹ and Alk² independently represent divalent C₁-C₃ alkylene radicals;

X represents -O-, -S-, -S(O)-, -S(O₂)-, -C(=O)-, -NH-, -NR₇- where R_7 is C_1 - C_3 alkyl;

and wherein

Alk¹ and Alk² and Z when other than hydrogen, independently are optionally substituted by

(C₁-C₃)alkyl, (C₂-C₃)alkenyl, or (C₂-C₃)alkynyl,

phenyl, optionally substituted by (C_1-C_3) alkyl, (C_1-C_3) alkoxy, halo, nitro, amino, mono- or di- (C_1-C_3) alkylamino, cyano or trifluoromethyl;

monocyclic 5 or 6-membered heterocyclic, optionally substituted by (C_1-C_3) alkyl, (C_1-C_3) alkoxy, halo, nitro, amino, mono- or di- (C_1-C_3) alkylamino, cyano or trifluoromethyl

benzyl, optionally substituted in the phenyl ring by (C_1-C_3) alkyl, (C_1-C_3) alkoxy, halo, nitro, amino, mono- or di- (C_1-C_3) alkylamino, cyano or trifluoromethyl,

hydroxy, phenoxy, (C_1-C_6) alkoxy, or hydroxyl (C_1-C_6) alkyl,

mercapto, (C₁-C₆) alkylthio or mercapto (C₁-C₆)alkyl,

oxo,

nitro,

cyano

halo

-COOH, or -COORA,

-COONH₂, -CONHR^A, or -CONR^AR^B

-CORA, -SO₂RA,

-NHCORA,

-NH₂, -NHR^A, or -NR^AR^B,

wherein R^A and R^B are independently a (C_1-C_6) alkyl group, R^A and R^B taken together with the nitrogen atom to which they are attached form a 5- or 6-membered heterocyclic ring which may be sutstituted by (C_1-C_3) alkyl, hydroxyl, or hydroxyl (C_1-C_3) alkyl.

15. (Currently Amended) A compound as claimed in any of claims claim 1 to 13 wherein A is a group -NR₈R₉ wherein R₈ and R₉ when taken together with the nitrogen atom to which they are attached form a saturated heterocyclic ring of 5 to 8 atoms optionally fused to a saturated or unsaturated carbocyclic or hetercyclic second ring of 5 to 7 ring atoms, any of which rings being optionally substituted by a radical of formula (IV)(II) as defined above

$$\frac{1}{2} (Alk^1)_m - (X)_p - (Alk^2)_n - Z \qquad (IV)$$

wherein

m, p and n are independently 0 or 1;

Z represents hydrogen or a carbocyclic or heterocyclic ring of 5 to 7 ring atoms which is optionally fused to a saturated or unsaturated carbocyclic or heterocyclic second ring of 5 to 7 ring atoms;

Alk¹ and Alk² independently represent divalent C₁-C₃ alkylene radicals;

X represents -O-, -S-, -S(O)-, -S(O₂)-, -C(=O)-, -NH-, -NR₇- where R_7 is C_1 - C_3 alkyl;

and wherein

Alk¹ and Alk² and Z when other than hydrogen, independently are optionally substituted by

 (C_1-C_3) alkyl, (C_2-C_3) alkenyl, or (C_2-C_3) alkynyl,

phenyl, optionally substituted by (C_1-C_3) alkyl, (C_1-C_3) alkoxy, halo, nitro, amino, mono- or di- (C_1-C_3) alkylamino, cyano or trifluoromethyl;

monocyclic 5 or 6-membered heterocyclic, optionally substituted by(C₁-C₃)alkyl, (C₁-C₃)alkoxy, halo, nitro, amino, mono- or di-(C₁-C₃)alkylamino, cyano or trifluoromethyl

benzyl, optionally substituted in the phenyl ring by (C_1-C_3) alkyl, (C_1-C_3) alkoxy, halo, nitro, amino, mono- or di- (C_1-C_3) alkylamino, cyano or trifluoromethyl,

hydroxy, phenoxy, (C_1-C_6) alkoxy, or hydroxyl (C_1-C_6) alkyl,

mercapto, (C₁-C₆) alkylthio or mercapto (C₁-C₆)alkyl,

oxo,

nitro,

cyano

halo

-COOH, or -COORA,

-COONH2, -CONHRA, or -CONRARB

 $-COR^A$, $-SO_2R^A$,

-NHCOR^A,

-NH₂, -NHR^A, or -NR^AR^B,

wherein R^A and R^B are independently a (C_1-C_6) alkyl group, R^A and R^B taken together with the nitrogen atom to which they are attached form a 5- or 6- membered heterocyclic ring which may be sutstituted by (C_1-C_3) alkyl, hydroxyl, or hydroxyl (C_1-C_3) alkyl.

16. (Original) A compound as claimed in claim 15 wherein R₈ and R₉ when taken together with the nitrogen atom to which they are attached form an optionally substituted 1-pyrrolidinyl, piperidin-1-yl, 1-piperazinyl, hexahydro-1-pyridazinyl, morpholin-4-yl, tetrahydro-1,4-thiazin-4-yl 1-oxide, tetrahydro-1,4-thiazin-4-yl, 1,1-dioxide, hexahydroazipino, thiomorpholino, diazepino, or thiazolidinyl ring.

17. (Original) A compound as claimed in claim 15 wherein R₈ and R₉ when taken together with the nitrogen atom to which they are attached form an optionally substituted piperidin-1-yl or 1-piperazinyl ring.

18. (Original) A compound as claimed in claim 14 wherein in the group (IV) Alk¹ and Alk² independently represent –(CH₂)- or –(CH₂CH₂)-.

19. (Currently Amended) A compound as claimed in claim 14 or claim 18—wherein in the group (IV) m is 0, p is 1, and X is -C(=0)- or $-S(O_2)$ -.

20. (Currently Amended) A compound as claimed in claim 1 of formulae (IIA) – (IID).

R¹ R²N-represents a cyclic amino group;

X₁ is a bond, C₁-C₃ alkylene, -NH- or-O-; and

A₁ is optionally substituted C₁-C₃ alkyl, cycloalkyl, aryl, or heterocyclic, for example methyl, ethyl phenyl, cyclopentyl, cyclohexyl, 2- or 3-furanyl, 2- or 3-thienyl, 2-, 3- or 4-pyridyl, 3-, 4- or 5-pyrazolyl, 3-, 4- or 5-oxazolyl, or 3-, 4- or 5-thiazolyl, methoxymethyl, 3,5-bis-(trifluoromethyl)phenyl, 4-trifluoromethylphenyl, 4-methoxyphenyl, 3,4-methylenedioxyphenyl, 4-fluorophenyl benzyl, 3-pyridyl, 4-pyridyl, cyclohexyl, 1,3-dimethylpyrazol-5-yl, 1-methylimidazol-5-yl, and 2-[morpholin-1-yl]pyrid-5-yl.

21. (Original) A compound as claimed in claim 20 wherein R₂ is n-propyl, n-butyl, n-pentyl, cyclopentylmethyl, cyclopentylmethyl or cyclohexylethyl;

X₁ is a bond, -CH₂-, -CH₂CH₂-, -CH₂CH₂CH₂-, -NH- or -O-; and

A₁ is methyl, ethyl phenyl, cyclopentyl, cyclohexyl, 2- or 3-furanyl, 2- or 3-thienyl, 2-, 3-or 4-pyridyl, 3-, 4- or 5-pyrazolyl, 3-, 4-or 5-oxazolyl, or 3-, 4-or 5-thiazolyl, methoxymethyl, 3,5-bis-(trifluoromethyl) phenyl, 4-trifluoromethylphenyl, 4-methoxyphenyl, 3,4-methylenedioxyphenyl, 4-fluorophenyl benzyl, 3-pyridyl, 4-pyridyl, cyclohexyl, 1,3-dimethylpyrazol-5-yl, 1-methylimidazol-5-yl, or 2-[morpholin-1-yl]pyrid-5-yl.

- 22. (Currently Amended) A method for the treatment of bacterial infections in humans and non-human mammals, which comprises administering to a subject suffering such infection an antibacterially effective dose of a compound as claimed in any of the preceding claims claim 1.
- 23. (Currently Amended) A method of inhibiting bacterial growth in vitro and in vivo in mammals comprising applying The use of a compound as claimed in claim 1 any of claims 1 to 21 for inhibiting bacterial growth in vitro and in vivo in mammals.
- 24. (Canceled)
- 25. (Currently Amended) A method for the treatment of bacterial contamination by applying an antibacterially effective amount of a compound as claimed in any of claims 1 to 21 claim 1

to the site of contamination.

26. (Currently Amended) A pharmaceutical or veterinary composition comprising a compound as claimed in any of claims 1 to 21 claim 1 together with a pharmaceutically or veterinarily acceptable carrier or excipient.